

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: Huiping GUAN et al.
Title: 3-(4-AMIDOPYRROL-2-YLMETHYLIDENE)-2-INDOLINONE
DERIVATIVES AS PROTEIN KINASE INHIBITORS
Prior Appl. No.: 10/076,140
Prior Appl. Filing Date: 02/15/2002
Examiner: Unassigned
Art Unit: Unassigned

INFORMATION DISCLOSURE STATEMENT
UNDER 37 CFR §1.56

Mail Stop PATENT APPLICATION
Commissioner for Patents
PO Box 1450
Alexandria, Virginia 22313-1450

Sir:

Applicants submit herewith on Form PTO/SB/08 a listing of the documents cited by or submitted to the U.S. PTO in parent application Serial No. 10/076,140, filed 02/15/2002. As provided in 37 CFR §1.98(d), copies of the documents are not being provided since they were previously submitted to the United States Patent & Trademark Office in the above-identified parent application.

The submission of any document herewith, which is not a statutory bar, is not intended as an admission that such document constitutes prior art against the claims of the present application or that such document is considered material to patentability as defined in 37 CFR §1.56(b). Applicants do not waive any rights to take any action which would be appropriate to antedate or otherwise remove as a competent reference any document which is determined to be a *prima facie* art reference against the claims of the present application.

TIMING OF THE DISCLOSURE

The listed documents are being submitted in compliance with 37 CFR §1.97(b), within three (3) months of the filing date of the application.

RELEVANCE OF EACH DOCUMENT

Applicants respectfully request that the listed documents be considered by the Examiner and be made of record in the present application and that an initialed copy of Form PTO/SB/08 be returned in accordance with MPEP §609.

The Commissioner is hereby authorized to charge any additional fees which may be required regarding this application under 37 CFR §§ 1.16-1.17, or credit any overpayment, to Deposit Account No. 19-0741. Should no proper payment be enclosed herewith, as by a check being in the wrong amount, unsigned, post-dated, otherwise improper or informal or even entirely missing, the Commissioner is authorized to charge the unpaid amount to Deposit Account No. 19-0741.

Date

9/8/03

FOLEY & LARDNER
Customer Number: 22428



22428

PATENT TRADEMARK OFFICE

Telephone: (202) 672-5475
Facsimile: (202) 672-5399

Respectfully submitted,

By

Beth A. Burrous
Attorney for Applicant
Registration No. 35,087

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Substitute for form 1449B/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT Date Submitted: (use as many sheets as necessary)				Complete if Known	
				Prior Application Number 10/076,140	
				Prior Appl. Filing Date February 15, 2002	
				First Named Inventor Huiping GUAN et al.	
				Group Art Unit Unassigned	
				Examiner Name Unassigned	
				Attorney Docket Number 034536-0188	
Sheet	1	of	30		

U.S. PATENT DOCUMENTS						
Examiner Initials*	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code ² (if known)			
	B1	Re. 35,096	E	Taniguchi et al.	11-21-1995	
	B2	Re. 36,256	E	Spada et al.	07-20-1999	
	B3	2,622,980		Copeland	12-23-1952	
	B4	2,872,372		Hull	02-03-1959	
	B5	2,968,557		Burgandt et al.	01-17-1961	
	B6	3,140,180		Fritz	07-07-1964	
	B7	3,308,134		Plostneiks	03-07-1967	
	B8	3,551,571		Pachter et al.	12-29-1970	
	B9	3,564,016		Schoen et al.	02-16-1971	
	B10	3,715,364		Hoff	02-06-1973	
	B11	3,880,871		Haugwitz et al.	04-29-1975	
	B12	3,922,163		Church et al.	11-25-1975	
	B13	4,002,643		Carson	01-11-1977	
	B14	4,002,749		Rovnyak	01-11-1977	
	B15	4,053,613		Rovnyak et al.	10-11-1977	
	B16	4,070,366		Gregorovich et al.	01-24-1978	
	B17	4,259,345		Cross et al.	03-31-1981	
	B18	4,343,923		Lenox et al.	08-10-1982	

Examiner Signature	Date Considered
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⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

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		Number	Kind Code ² (if known)			
	B19	4,376,110		David et al.	03-08-1983	
	B20	4,436,892		Zondler et al.	03-13-1984	
	B21	4,489,089		Wright, Jr. et al.	12-18-1984	
	B22	4,493,842		Furazawa et al.	01-15-1985	
	B23	4,560,700		Schnettler et al.	12-24-1985	
	B24	4,628,105		Schmid et al.	12-09-1986	
	B25	4,642,309		Michel et al.	02-10-1987	
	B26	4,678,798		Rentzea et al.	07-07-1987	
	B27	4,826,847		Michel et al.	05-02-1989	
	B28	4,853,403		Shiraishi et al.	08-01-1989	
	B29	4,853,404		Takamura et al.	08-01-1989	
	B30	4,868,304		Larock	09-19-1989	
	B31	4,924,000		Rentzea et al.	05-08-1990	
	B32	4,966,849		Vallee et al.	10-30-1990	
	B33	4,971,996		Shiraishi et al.	11-20-1990	
	B34	4,987,146		Rohde et al.	01-22-1991	
	B35	5,043,348		Zoller et al.	08-27-1991	
	B36	5,043,454		Wriede et al.	08-27-1991	

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	B37	5,047,554		Ehrgott et al.	09-10-1991	
	B38	5,051,417		Nadler et al.	09-24-1991	
	B39	5,057,538		Shiraishi et al.	10-15-1991	
	B40	5,082,856	A	Taniguchi et al.	01-21-1992	
	B41	5,089,516	A	Shiraishi et al.	02-18-1992	
	B42	5,124,347	A	Connor et al.	06-23-1992	
	B43	5,145,983	A	West	09-08-1992	
	B44	5,153,217	A	Taniguchi et al.	10-06-1992	
	B45	5,196,446	A	Levitzi et al.	03-23-1993	
	B46	5,202,341	A	Shiraishi et al.	04-13-1993	
	B47	5,206,261	A	Kawaguchi et al.	04-27-1993	
	B48	5,217,999	A	Levitzi et al.	06-08-1993	
	B49	5,258,357	A	Muenster et al.	11-02-1993	
	B50	5,278,184	A	Artico et al.	01-11-1994	
	B51	5,290,947	A	Zoller et al.	03-01-1994	
	B52	5,302,606	A	Spada et al.	04-12-1994	
	B53	5,322,950	A	Sircar et al.	06-21-1994	
	B54	5,330,992	A	Eissenstat et al.	07-19-1994	

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	B55	5,332,736	A	Carmosin et al.	07-26-1994	
	B56	5,374,652	A	Buzzetti et al.	12-20-1994	
	B57	5,382,593	A	Le Baut et al.	01-17-1995	
	B58	5,389,661	A	Sircar et al.	02-14-1995	
	B59	5,397,787	A	Buzzetti et al.	03-14-1995	
	B60	5,409,930	A	Spada et al.	04-25-1995	
	B61	5,409,949	A	Buzzetti et al.	04-25-1995	
	B62	5,463,052	A	Haga et al.	10-31-1995	
	B63	5,565,324	A	Still et al.	10-15-1996	
	B64	5,610,173	A	Schwartz et al.	03-11-1997	
	B65	5,723,665	A	Kato et al.	03-03-1998	
	B66	5,786,488	A	Tang et al.	07-28-1998	
	B67	5,792,783	A	Tang et al.	08-11-1998	
	B68	5,834,504	A	Tang et al.	11-10-1998	
	B69	5,849,710	A	Battistini et al.	12-15-1998	
	B70	5,880,141	A	Tang et al.	03-09-1999	
	B71	5,883,113	A	Tang et al.	03-16-1999	
	B72	5,883,116	A	Tang et al.	03-16-1999	

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		Number	Kind Code ² (if known)			
	B73	5,886,020	A	Tang et al.	03-23-1999	
	B74	6,130,239	A	Chen et al.	10-10-2000	
	B75	6,133,305	A	Tang et al.	10-17-2000	
	B76	6,284,894	B1	Phillion et al.	09-04-2001	
	B77	6,310,217	B1	Lehr	10-30-2001	
	B78	6,395,736		Parks et al.	05-28-2002	
	B79	6,451,838		Moon et al.	09-17-2002	
	B80	6,462,072		Hamilton et al.	10-08-2002	

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		Office ³	Number ⁴	Kind Code ⁵ (if known)				
	B81	WO	88/07035	A1	KANEGAFUCHI KAGAKU KOGYO KABUSHIKI KAISHA	09-22-1988		
	B82	WO	91/13055	A2	FARMITALIA CARLO ERBA SRL	09-05-1991		
	B83	WO	91/15495	A1	PFIZER INC.	10-17-1991		
	B84	WO	92/03736	A1	SEIKAGAKU KOGYO KABUSHIKI KAISHA	03-05-1992		
	B85	WO	92/07830	A2	PFIZER INC.	05-14-1992		
	B86	WO	92/20642	A1	RHONEPOULENC RORER INTERNATIONAL	11-26-1992		
	B87	WO	92/21660	A1	PFIZER INC.	12-10-1992		

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	B88	WO	93/01182	A1	FARMITALIA CARLO ERA SRL	01-21-1993		
	B89	WO	93/23040	A1	MERCK & CO., INC.	11-25-1993		
	B90	WO	94/03427	A1	WARNER-LAMBERT COMPANY	02-17-1994		
	B91	WO	94/10202	A1	GENENTECH, INC.	05-11-1994		
	B92	WO	94/14808	A1	FARMITALIA CARLO ERBA SRL	07-07-1994		
	B93	WO	95/01349	A1	FARMITALIA CARLO ERBA SRL	01-12-1995		
	B94	WO	95/14667	A1	PFIZER INC.	06-01-1995		
	B95	WO	95/17181	A1	PHARMACIA S.P.A.	06-29-1995		
	B96	WO	95/24190	A2	SUGEN, INC.	09-14-1995		
	B97	WO	96/00226	A1	PHARMACIA S.P.A.	01-04-1996		
	B98	WO	96/16964	A1	PHARMACIA S.P.A.	06-06-1996		
	B99	WO	96/22976	A1	PHARMACIA S.P.A.	08-01-1996		
	B100	WO	96/32380	A1	PHARMACIA S.P.A.	10-17-1996		
	B101	WO	96/40116	A1	SUGEN, INC.	12-19-1996		
	B102	WO	97/25986	A1	TAIHO PHARMACEUTICAL CO., LTD.	07-24-1997		
	B103	WO	97/34920	A1	SUGEN, INC.	09-25-1997		
	B104	WO	97/36867	A1	PFIZER, INC.	10-09-1997		
	B105	WO	98/07695	A1	SUGEN, INC.	02-26-1998		

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	B106	WO	98/07835	A2	SUGEN, INC.	02-26-1998		
	B107	WO	98/24432	A2	SUGEN, INC.	06-11-1998		
	B108	WO	98/38984	A2	SUGEN, INC.	09-11-1998		
	B109	WO	98/45708	A1	SUGEN, INC.	10-15-1998		
	B110	WO	98/50356	A1	SUGEN, INC.	11-12-1998		
	B111	WO	98/56376	A1	SUGEN, INC.	12-17-1998		
	B112	WO	99/10325	A1	GLAXO GROUP LIMITED	03-04-1999		
	B113	WO	99/19325	A1	SYNTHELABO	04-22-1999		
	B114	WO	99/48868	A2	SUGEN, INC.	09-30-1999		
	B115	WO	99/52869	A1	BOEHRINGER INGELHEIM PHARMA KG	10-21-1999		
	B116	WO	99/65869	A1	BAYER AKTIENGESELLSCHAFT	12-23-1999		
	B117	WO	99/61422	A1	SUGEN, INC.	12-02-1999		
	B118	WO	00/35920	A2	F. HOFFMANN-LA ROCHE AG	06-22-2000		
	B119	WO	00/38519	A1	SUGEN, INC.	07-06-2000		
	B120	WO	00/08202	A2	SUGEN, INC.	02-17-2000		
	B121	WO	00/56709	A1	SUGEN, INC.	09-28-2000		
	B122	WO	01/60814	A2	SUGEN, INC.	08-23-2001		
	B123	WO	01/90068	A2	SUGEN, INC. et al.	11-29-2001		
	B124	DE	878,539		Von FREYBERG, et al.	06-05-1953		X

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				Prior Application Number		10/076,140
				Prior Appl. Filing Date		February 15, 2002
				First Named Inventor		Huiping GUAN et al.
				Group Art Unit		Unassigned
				Examiner Name		Unassigned
Sheet 8 of 30				Attorney Docket Number		034536-0188

FOREIGN PATENT DOCUMENTS								
Examiner Initials*	Cite No. ¹	Foreign Patent Document			Name of Patentee or Applicant of Cited Documents	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁶
		Office ³	Number ⁴	Kind Code ⁵ (if known)				
	B125	DE	2,159,360	A	BAYER AG	06-14-1973		X
	B126	DE	2,159,361	A	BAYER AG	06-14-1973		X
	B127	DE	2,159,362		BAYER AG	06-14-1973		
	B128	DE	2,159,363	A	BAYER AG	06-14-1973		X
	B129	DE	2,321,656	A	COLGATE-PALMOLIVE CO.	11-15-1973		X
	B130	DE	3,426,419	A	BOEHRINGER MANNHEIM GMBH	01-23-1986		X
	B131	EP	0 252 713	B1	PFIZER INC.	01-13-1988		
	B132	EP	0 304 493	B1	KANEGAFUCHI KAGAKU KOGTO KABUSHIKI KAISHA	03-01-1989		
	B133	EP	0 351 213	A2	LES LABORATOIRES BEECHAM S.A.	01-17-1990		
	B134	EP	0 525 472	A2	FARMITALIA CARLO ERBA SRL	02-03-1993		
	B135	EP	0 566 226	B1	ZENECA LIMITED	10-20-1993		
	B136	EP	0 580 502	B1	ADIR ET COMPAGNIE	01-26-1994		X
	B137	EP	0 626 377	B1	SHIONOGI & CO., LTD.	11-30-1994		
	B138	EP	0 632 102	A1	BAYER AG	01-04-1995		X
	B139	EP	0 662 473	A1	PHARMACIA S.P.A.	07-12-1995		
	B140	EP	0 788 890	A1	AGFA-GEVAERT	08-13-1997		
	B141	EP	0 810 217	A1	JAPAN ENERGY CORPORATION	12-03-1997		
	B142	EP	0 769 947	B1	TANG, Peng Cho et al.	05-02-1997		

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	B143	EP	0 934 931	A2	SUGEN, INC.	08-11-1999		
	B144	EP	1 082 305	A1	SUGEN, INC.	03-14-2001		
	B145	FR	1.398.224		IMPERIAL CHEMICAL INDUSTRIES LIMITED	05-07-1965		X
	B146	FR	1.599.772		INSTITUT PASTEUR	08-28-1970		X
	B147	FR	2.689.397	A1	ADIR ET COMPAGNIE	10-08-1993		X
	B148	GB	809,691		Roy HULL	03-04-1959		
	B149	GB	835,473		Norman SENIOR	05-18-1960		
	B150	JP	62-29570	A	KANEGAFUCHI CHEM KK	02-07-1987		X
	B151	JP	62-39564	A	KANEGAFUCHI CHEM KK	02-20-1987		X
	B152	JP	63-141955	A	KANEGAFUCHI CHEM KK	06-14-1988		X
	B153	JP	5-58894	A	KANEKA CORP	03-09-1993		X
	B154	CA	2,012,634	A1	UNIVERSITY OF BRITISH COLUMBIA	09-20-1991		
	B155	AU	286870		IMPERIAL CHEMICAL INDUSTRIES OF AUSTRALIA AND NEW ZEALAND LIMITED	05-11-1967		

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				Examiner Name	Unassigned
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	B156	ABRAMOVITCH and HEY, "Internuclear cyclisation. Part VIII. Naphth[3:2:1-cd]oxindoles," <u>J. Chem. Soc.</u> 1697-1703 (1954), Strand, London		
	B157	ABRAMOVITCH et al., "A Novel Synthesis of a Cyclic Hydroxamic Acid Involving a Molecular Rearrangement," <u>Chemistry and Industry</u> 44:1871 (1967) 8Laporte Industries Limited, Lancashire		
	B158	Beilstein Reg. No. 236050, Beilstein Reference No. 4-21-00-06355		
	B159	AKBASAK and SUNAR-AKBASAK, "Oncogenes: cause or consequence in the development of glial tumors," <u>J. Neurol. Sci.</u> 111:119-133 (1992)8 Elsevier Science Publishers		
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	B162	ANDREANI et al., "Synthesis and cardiotoxic activity of 2-indolinones bearing pyridyl groups," <u>Eur. J. Med. Chem.</u> 28:653-657 (1993) 8 Elsevier, Paris		
	B163	ANDREANI et al., "Synthesis and cardiotoxic activity of 2-indolinones," <u>Chemical Abstracts</u> , Vol. 113, abstract no. 78106 (1990)		
	B164	ANDREANI et al., "Synthesis and cardiotoxic activity of pyridylmethylene-2-indolinones," <u>Eur. J. Med. Chem.</u> 27:167-170 (1992) 8 Elsevier, Paris		
	B165	ANDREANI et al., "Synthesis and potential coanthracyclic activity of substituted 3-(5-imidazo[2,1-b]thiazolymethylene)-2-indolinones," <u>Eur. J. Med. Chem.</u> 32:919-924 (1997)8 Elsevier, Paris		
	B166	ANDREANI et al., "Synthesis of lactams with potential cardiotoxic activity," <u>Eur. J. Med. Chem.</u> 28:825-829 (1993)		
	B167	ANDREANI et al., "In Vivo Cardiotoxic Activity of Pyridylmethylene-2-indolinones," <u>Arzneimittel-Forschung Drug Research</u> 48:727-729 (1998) 8		

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	B168	ARTEAGA et al., "Blockade of the Type I Somatomedin Receptor Inhibits Growth of Human Breast Cancer Cells in Athymic Mice," <u>J. Clin. Invest.</u> 84:1418-1423 (1989) copyright The American Society for Clinical investigation, Inc.		
	B169	ARVIDSSON et al., "Tyr-716 in the Platelet-Derived Growth Factor β -Receptor Kinase Insert is Involved in GRB2 Binding and Ras Activation," <u>Molecular and Cellular Biology</u> 14:6715-6726 (1994) 8 The American Society for Microbiology		
	B170	AUTREY and TAHK, "The Synthesis and Stereochemistry of Some Isatylideneacetic Acid Derivatives," <u>Tetrahedron</u> 23:901-917 (1967) 8 Pergamon Press		
	B171	BAHNER and BROTHERTON, "6-Dimethylaminochrysene and Other Analogs of 4-(4-Dimethylamino)stilbene," <u>J. Med. Chem.</u> 12:722-723 (1969)		
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	B173	BAMFIELD et al., "Diels-Alder Reactions of Oxindolylideneacetone," <u>J. Chem. Soc. (C)</u> 1028-1030 (1966) 8		
	B174	BARBIER, et al., "Synthesis of Isobrassilexin, A Biologically Active Isomer of Brassilexin, a Cruciferae Phytoalexin," <u>Synthetic Communications</u> 23(22):3109-3117 (1993) 8 Marcel Dekker, Inc.		
	B175	BASERGA, "Oncogenes and the Strategy of Growth Factors," <u>Cell</u> 79:927-930 (1994) 8 Cell Press		
	B176	BASERGA, "The Insulin-like Growth Factor I Receptor: A Key to Tumor Growth?" <u>Cancer Research</u> 55:249-252 (1995)		

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	B177	Beilstein Reg. No. 233511 (1997)	
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	B179	Beilstein Reg. No. 252929 (1998)	
	B180	BENZIES, et al., "2-Formyl-3-Methoxymethylindole, 3-Ethoxymethyl-2-Formylindoline and 2-Formyl-3-Methylindole," <u>Synthetic Communications</u> : 16(14), 1799-1807 (1986) 8 Mercel Dekker, Inc.	
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	B184	BONNER et al., "Structure and Biological Activity of Human Homologs of the <i>raf/mil</i> Oncogene," <u>Molecular and Cellular Biology</u> 5:1400-1407 (1985) 8 The American Society for Microbiology	
	B185	BORSCHKE et al., "Über vielkernige kondensierte Systeme mit heterocyclischen Ringen. XIII.," <u>Liebigs Ann. Chem.</u> 550:160-174 (1941)	

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	B186	BUZZETTI et al., "Cinnamamide Analogs as Inhibitors of Protein Tyrosine Kinases," <u>Il Farmaco</u> 48:615-636 (1993)		
	B187	CANCE et al., "Novel Protein Kinases Expressed in Human Breast Cancer," <u>Int. J. Cancer</u> 54:571-577 (1993) 8 Wiley-Liss, Inc.		
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	B195	CODA et al., "(Z)- and (E)-Arylidene-1,3-dihydroindol-2-ones: Configuration, Conformation and Infrared Carbonyl Stretching Frequencies," <u>J. Chem. Soc. Perkin Trans. II</u> : 615-619 (1984)		
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				First Named Inventor	Huiping GUAN et al.
				Group Art Unit	Unassigned
				Examiner Name	Unassigned
Date Submitted:				Attorney Docket Number	034536-0188
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	B219	FOLKMAN, "Angiogenesis in Psoriasis: Therapeutic Implications," <u>J. Invest. Dermatol.</u> 59:40-43 (1973) copyright The Williams & Wilkins Co.		
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	B228	HODGES et al., "Chemical and biological properties of some oxindolidyl-3-methines," <u>Canadian J. Chemistry</u> 46:2189-2194 (1968)		
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	B267	MORETO et al., "Study of the Laxative Properties of the Disodium Salt of the Sulfuric Diester of 3,3 Bis-(4-Hydroxyphenyl)-7-Methyl-2-Indolinone (DAN-603) in the Rat," <u>European Journal of Pharmacology</u> 36:221-226 (1976) 8North-Holland Publishing Company		
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	B270	MOSMANN, "Rapid Colorimetric Assay for Cellular Growth and Survival: Application to Proliferation and Cytotoxicity Assays," <u>J. Immunol. Methods</u> 65:55-63 (1983) copyright Elsevier Publishers B.V.		
	B271	NEBER and RÖCKER, "On the action of benzaldehydes on the free o-aminophenylacetic acid (II)," <u>Chem. Ber.</u> 56:1710-1716 (1923) (GERMAN AND ENGLISH TRANSLATION)		
	B272	NISHIMURA et al., "Two Signaling Molecules Share a Phosphotyrosine-Containing Binding Site in the Platelet-Derived Growth Factor Receptor," <u>Molecular and Cellular Biology</u> 13:6889-6896 (1993)		
	B273	NODIFF et al., "Antimalarial Phenanthrene Amino Alcohols. 1. Fluorine-Containing 3- and 6-Substituted 9-Phenanthrenemethanols," <u>J. Med. Chem.</u> 14:921-925 (1971)		
	B274	NODIFF et al., "Antimalarial Phenanthrene Amino Alcohols. 3. Halogen-containing 9-phenanthrenemethanols," <u>Chemical Abstracts</u> , Vol. 83, abstract no. 188214 (1975)		
	B275	OSBORNE et al., "Effect of Estrogens and Antiestrogens on Growth of Human Breast Cancer Cells in Athymic Nude Mice," <u>Cancer Research</u> 45:584-590 (1985)		

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				Prior Appl. Filing Date	February 15, 2002
				First Named Inventor	Huiping GUAN et al.
				Group Art Unit	Unassigned
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	B276	O'SULLIVAN and ROTHERY, "The Preparation and Possible Clinical Significance of 4'-Dialkylaminoisoindogenides," <u>Clinica Chimica Acta</u> 62:181-182 (1975) 8Elsevier Scientific Publishing Company	
	B277	OZZELLO and SORDAT, "Behavior of Tumors Produced by Transplantation of Human Mammary Cell Lines in Athymic Nude Mice," <u>Eur. J. Cancer</u> 16:553-559 (1980)	
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	B279	PERKIN et al., "Harmine and Harmaline. Part II. The Synthesis of isoHarman," <u>J. Chem. Soc.</u> 103:1973-1985 (1913)	
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	B281	PLOWMAN et al., "Receptor Tyrosine Kinases as Targets for Drug Intervention," <u>DN&P</u> 7:334-339 (1994)	
	B282	QUALLICH et al., A General Oxindole Synthesis," <u>J. Synthetic Organic Chemistry</u> : 51-51 (1993)	
	B283	QUINN et al., "Fetal liver kinase 1 is a receptor for vascular endothelial growth factor and is selectively expressed in vascular endothelium," <u>Proc. Natl. Acad. Sci. USA</u> 90:7533-7537 (1993)	
	B284	ROZAKIS-ADCOCK et al., "Association of the Shc and Grb2/Sem5 SH2-containing proteins is implicated in activation of the Ras pathway by tyrosine kinases," <u>Nature</u> 360:689-692 (1992)	

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	B285	RUVEDA and GONZALEZ, "Geometric isomerism in benzylideneoxindoles," <u>Spectrochimica Acta</u> 26A:1275-1277 (1970)		
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	B288	SAITO and STREULI, "Molecular Characterization of Protein Tyrosine Phosphatases," <u>Cell Growth & Differentiation</u> 2:59-65 (1991) 8Molecular Biology Journal of the American Association for Cancer Research		
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	B292	SCHUCHTER et al., "Successful Treatment of Murine Melanoma with Bryostatin 1," <u>Cancer Research</u> 51:682-687 (1991)		
	B293	SEIBERT et al., "Clonal Variation of MCF-7 Breast Cancer Cells <u>in Vitro</u> and in Athymic Nude Mice," <u>Cancer Research</u> 43:2223-2234 (1983)		

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	B294	SHAFIE and GRANTHAM, "Role of Hormones in the Growth and Regression of Human Breast Cancer Cells (MCF-7) Transplanted Into Athymic Nude Mice," <u>J. Natl. Cancer Institute</u> 67:51-56 (1981)		
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	B296	SHIRAISHI et al., "Specific inhibitors of Tyrosine-Specific Protein Kinase, Synthetic 4-Hydroxycinnamamide Derivatives," <u>Biochemical and Biophysical Research Communications</u> 147:322-328 (1987) Academic Press		
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	B298	SHWEIKI et al., "Vascular endothelial growth factor induced by hypoxia may mediate hypoxia-initiated angiogenesis," <u>Nature</u> 359:843-845 (1992)		
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	B301	SKEHAN et al., "New Colorimetric Cytotoxicity Assay for Anticancer-Drug Screening," <u>J. Natl. Cancer Inst.</u> 82:1107-1112 (1990)		
	B302	SLAMON et al., "Studies of the HER-2/neu Proto-oncogene in Human Breast and Ovarian Cancer," <u>Science</u> 244:707-712 (1989)		

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	B304	SONGYANG et al., "SH2 Domains Recognize Specific Phosphopeptide Sequences," <u>Cell</u> 72:767-778 (1993) 8 Cell Press	
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	B311	SUN et al., "Design, Synthesis, and Evaluations of Substituted 3-[(3- or 4-Carboxyethylpyrrol-2-yl) methylidenyl]indolin-2-ones as Inhibitors of VEGF, FGF, and PDGF Receptor Tyrosine Kinases," <u>Journal of Medicinal Chemistry</u> 42: 5120-5130 (1999) 8American Chemical Society	

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	B320	THOMPSON et al., "Facile Dimerisation of 3-Benzylideneindoline-2-thiones," <u>J. Chem. Soc. Perkin Trans. (I)</u> 1835-1837 (1993)		

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	B327	VAISMAN et al., "Characterization of the Receptors for Vascular Endothelial Growth Factor," <u>J. Biol. Chem.</u> 265:19461-19466 (1990) 8 The American Society for Biochemistry and Molecular Biology	
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	B330	WAHL et al., "3-benzilidene-5-methyl-1,3-dihydroindol-2-one," <u>Ann. Chim.</u> 350 (1926), DATABASE CROSSFIRE, Beilstein Reference No. 2-21-00-00290	
	B331	WAHL et al., "Chimie Organique - Sur les iso-indogenides," <u>C.R. Hebd. Seances Acad. Sci.</u> 149:132-134 (1909)	
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	B334	WALKER, "Synthesis of a α -(p-Aminophenyl)- and α -(p-Chlorophenyl)- β -aryl-propionitriles by Catalytic Reduction of Stilbenenitriles," <u>J. Med. Chem.</u> 8:583-588 (1965)	
	B335	WALKER et al., "Synthesis of New 3-(Pyridylmethylene)-, 3-(Pyridylmethyl)-, 3-(Piperidylmethyl)-, and 3-(β -Alkylaminoethyl)-2-indolinones. The Reduction of Isoindogenides, Nitro Compounds, and Pyridines in a Series of 2-Indolinones," <u>J. Med. Chem.</u> 8:626-637 (1965)	
	B336	WARRI et al., "Estrogen Suppression of erbB2 Expression is Associated with Increased Growth Rate of ZR-75-I Human Breast Cancer Cells <u>In Vitro</u> and in Nude Mice," <u>Int. J. Cancer</u> 49:616-623 (1991) 8 Wiley-Leiss, Inc.	
	B337	WEIDNER et al., "Tumor Angiogenesis and Metastasis -- Correlation in Invasive Breast Carcinoma," <u>New England J. Medicine</u> 324:1-7 (1991) 8 Massachusetts Medical Society	
	B338	WINKELMANN et al., "Chemotherapeutically Active Nitro Compounds: 4. 5-Nitroimidazoles (Part I)," <u>Arzneim.-Forsch./Drug Res.</u> 27:2251-2263 (1977)	

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT Date Submitted: <i>(use as many sheets as necessary)</i>				Prior Application Number	10/076,140
				Prior Appl. Filing Date	February 15, 2002
				First Named Inventor	Huiping GUAN et al.
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				Examiner Name	Unassigned
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	B339	WRIGHT et al., "Cyclic Hydroxamic Acids Derived from Indole," <u>J. Am. Chem. Soc.</u> 78:221-224 (1956)	
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	B342	ZAMAN et al., "Tyrosine Kinase Activity of Purified Recombinant Cytoplasmic Domain of Platelet-Derived Growth Factor β -Receptor (β -PDGFR) and Discovery of a Novel Inhibitor of Receptor Tyrosine Kinases," <u>Biochemical Pharmacology</u> 57:57-64 (1999) 8Elsevier Science Inc.	
	B343	ZHANG et al., "Microtubule Effects of Welwistatin, a Cyanobacterial Indolinone that Circumvents Multiple Drug Resistance," <u>Molecular Pharmacology</u> 49:228-234 (1996) 8The American Society for Pharmacology and Experimental Pharmaceutics	
	B344	ZHUNGIETU et al., "Reaction of Indoles and 2-Ketoindolines With Some Aldehydes," <u>Chemical Abstracts</u> , Vol. 78, abstract no. 111201 (1990)	

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